

## Enantiospecific Total Synthesis of the Hapalindoles, Fischerindoles, and Welwitindolinones *via* a Redox Economic Approach

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**Abstract:** Full details are provided for the total synthesis of several members of the hapalindole family of natural products, including hapalindole Q, 12-*epi*-hapalindole D, 12-*epi*-fischerindole U, 12-*epi*-fischerindole G, 12-*epi*-fischerindole I, and welwitindolinone A. Use of the recently developed direct indole coupling enabled an efficient, practical, scalable, and protecting-group-free synthesis of each of these natural products. The original biosynthetic proposal is reviewed, and a revised biosynthetic hypothesis is suggested, validated by the above syntheses. The syntheses are also characterized by an adherence to the concept of “redox economy”. Analogous to “atom economy” or “step economy”, “redox economy” minimizes the superfluous redox manipulations within a synthesis; rather, the oxidation state of intermediates linearly and steadily increases throughout the course of the synthesis.

### Introduction

The first members of the hapalindole-type natural products were isolated from the Stigonemataceae family of cyanobacteria in 1984 by Moore and colleagues.<sup>1</sup> In the 24 years since their initial discovery, 63 members<sup>2</sup> have been added to this family of alkaloids, which include the hapalindoles, fischerindoles, welwitindolinones, ambiguines, hapalindolinones, hapaloxindoles, and fontonamides (see Chart 1). These natural products have been isolated from soil samples in a myriad of habitats around the globe,<sup>3</sup> and a broad range of biological activities arises from the different structural classes. Insecticidal activity is observed for several hapalindoles<sup>21</sup> and welwitindolinones.<sup>2f</sup>

Reports have surfaced of anti-algal activity from the hapalindoles,<sup>1</sup> antimycotic activity from the hapalindoles,<sup>1,2e,i</sup> welwitindolinones,<sup>2f</sup> and ambiguines,<sup>2d,k</sup> and antibacterial activity from the hapalindoles<sup>2m,4</sup> and ambiguines.<sup>2k</sup> Additionally, it has been found that hapalindolinone A inhibits arginine vasopressin binding.<sup>2b</sup> Finally, potent anticancer activity against multiple drug-resistant ovarian cancer cell lines has been reported for the welwitindolinones,<sup>2f,5</sup> which apparently exert this effect through microtubule depletion.<sup>6</sup>

Not only do a vast number of the members of this natural product family exhibit potent and exciting biological activities, but they also all contain intriguing and unprecedented molecular architectures. Though distinct, they are united by several structural features, most notably an indole (or indole-derived) heterocycle with a monoterpene unit appended at C(3) (see Scheme 1 for numbering), comprising the core of these molecules. All but a handful of them contain an isonitrile or an isothiocyanate at C(11), with an all-carbon quaternary center, comprised of a methyl and vinyl group, vicinal to this moiety (C(12)). The hapalindoles are the simplest members of this family, containing the core structure described above, housed within a tricyclic framework. Many contain further functionalization, in the form of either unsaturation (at C(10)) or chlorination (at C(13)), and several contain an additional carbocycle arising from the union of C(4) of indole with the isopropylidene unit at C(15). The fischerindoles, tetracycles

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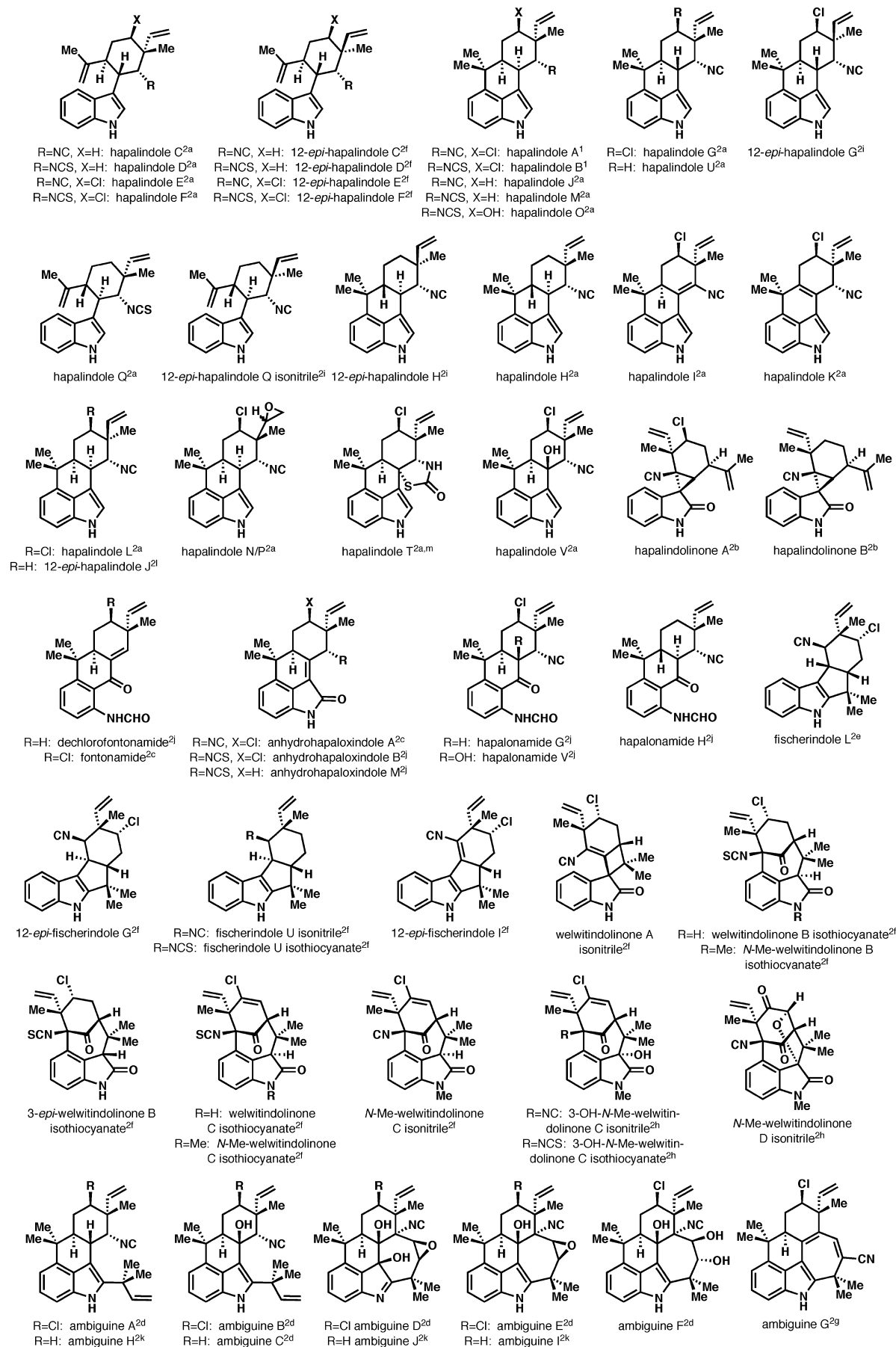
(3) These locations include the Marshall Islands (ref 1), the Everglades in Florida (ref 2b), Australia (ref 2f), Micronesia (ref 2h), Papua New Guinea (ref 2i), and Israel (ref 2k).

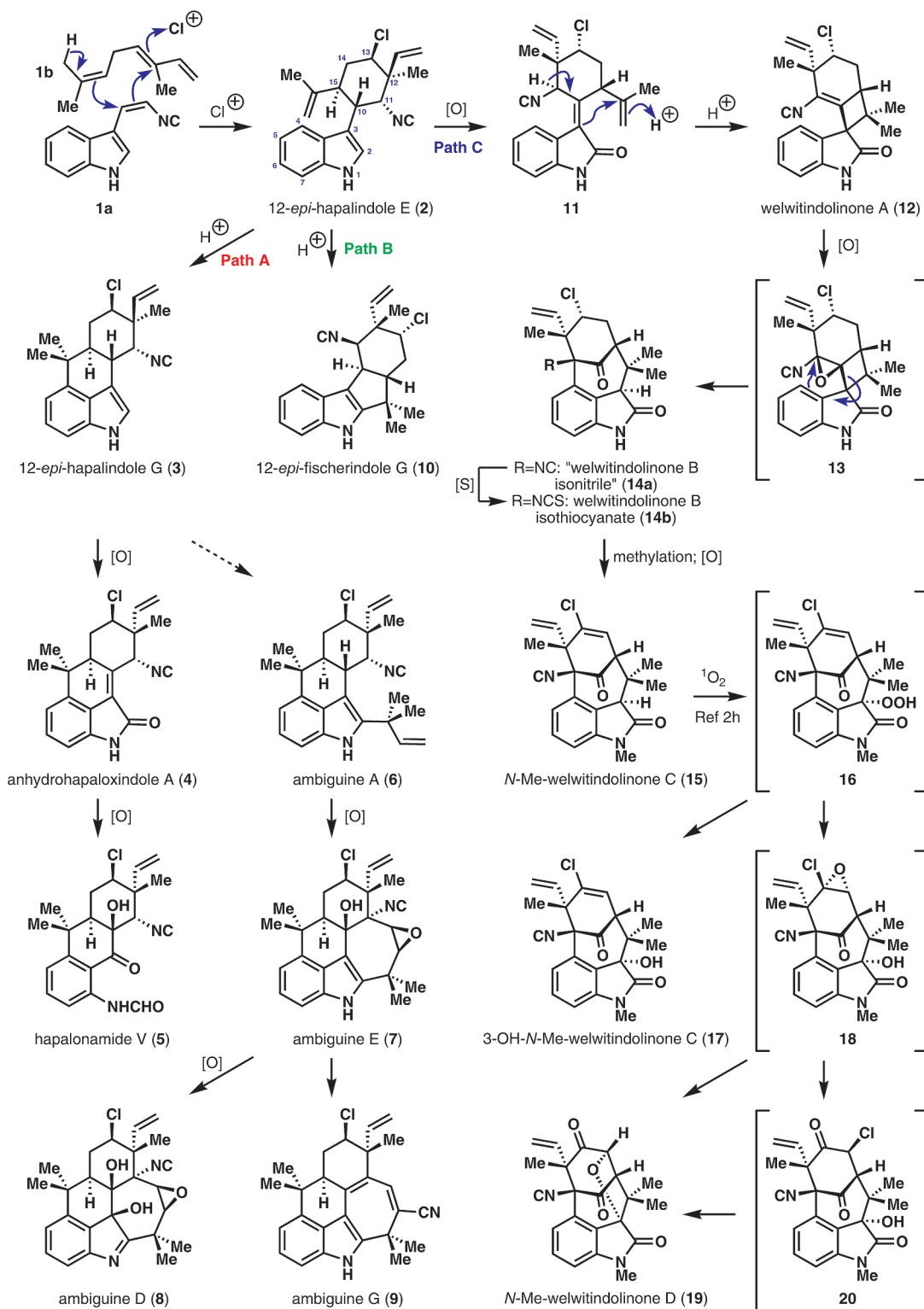
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Chart 1. All Known Hapalindole-Type Natural Products



Scheme 1. Moore's Proposed Biosynthetic Relationships between the Hapalindole-Type Alkaloids<sup>a</sup>

<sup>a</sup> [O], oxidation; [S], sulfur insertion.

formed *via* the union of C(2) of indole with the isopropylidene unit at C(15), are also characterized by varying degrees of functionalization and oxidation. The hapaloxindoles and fon-tonamides are structurally related to the tetracyclic hapalindoles, although the indole has been either oxidized to give the oxindole or oxidatively cleaved to form the formylkynurenine. Also related to the hapalindoles, albeit much more complex, are the ambiguines, which contain additional functionalization at C(2)

of indole, specifically a *tert*-prenyl moiety. In the most complex ambiguines, this *tert*-prenyl group is further cyclized and oxidized. In addition to the unifying structural features of this family of natural products, the hapalindolinones contain a unique component within their molecular architecture, specifically a spirocyclic cyclopropane that joins C(11) with C(3) of the oxindole heterocycle. Finally, the welwitindolinones are found in one of two structural classes, the first being welwitindolinone

A, which contains a spirocyclic cyclobutane centered around C(3). The remaining welwitindolinones are comprised of a [4.3.1]bicyclononanone core, which contains an assortment of oxidative functionalization.

With a family of such diverse and unique molecular architectures, it should come as no surprise that several syntheses have been reported for these natural products, specifically focusing on the simpler members of the family. Syntheses have been reported for hapalindoles G,<sup>7</sup> H,<sup>8</sup> J,<sup>9</sup> M,<sup>9</sup> Q,<sup>10</sup> O,<sup>11</sup> and U,<sup>8</sup> in addition to approaches to various other hapalindoles.<sup>12</sup> There are no reported efforts toward the hapalindolinones, hapaloxindoles and fontonamides, and only one approach toward an ambiguine.<sup>13</sup> The first total synthesis of an ambiguine (ambiguine H) was reported from our laboratory.<sup>14</sup> Despite the many approaches toward the welwitindolinones,<sup>15</sup> at the time of our initial communication<sup>16</sup> of the work presented herein, the members of the welwitindolinone family had not yet succumbed to synthesis; however, the Wood group reported a very elegant synthesis of welwitindolinone A shortly thereafter.<sup>17</sup>

### Biosynthetic Relationships and Retrosynthetic Analysis

Biomimetic syntheses are often more efficient due to the tactics that Nature employs, namely rapid assembly of skeletal complexity, a linear increase of oxidation state, use of mild and simple reagents, and the ability to control chemoselectivity (lack of protecting groups).<sup>18</sup> Despite their inherent advantages, biomimetic syntheses can be exceedingly difficult, due to the

inability of chemists to attain the chemo-, regio-, and stereo-control characterizing most enzymatic processes. The careful practitioner can make use of many abiotic tools in solving these problems; however, these methods usually demand significant departure from the ideal biomimetic route. In light of these difficulties, it is certainly possible, and perhaps prudent, to find an appropriate balance when designing a retrosynthesis. An ideal synthesis might entail the use of powerful synthetic methods, coupled with a flexible adherence to the general synthetic blueprint provided by Nature.

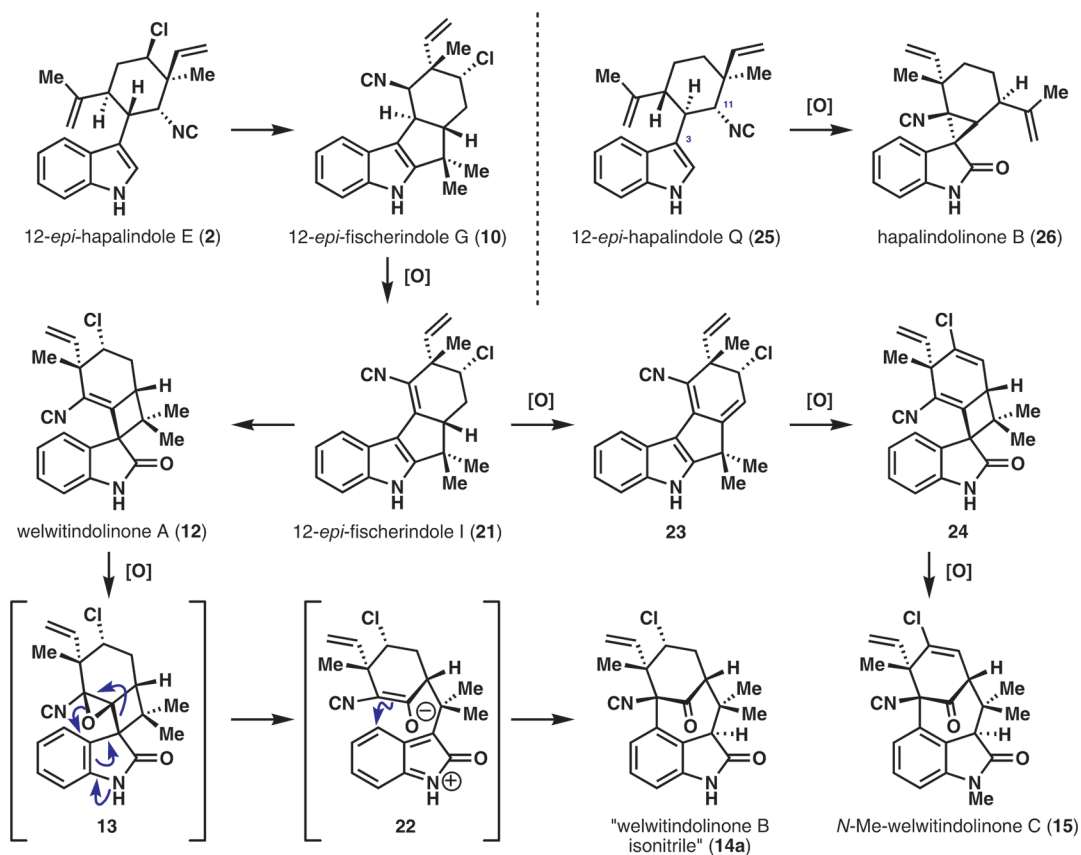
Given these considerations, and with such a large and diverse family of complex natural products, a biosynthetic proposal that comprehensively describes the interrelationships between the members would undoubtedly be enlightening to any synthetic undertaking. The Moore group, in conjunction with their elegant isolation studies, put forth many plausible biosynthetic ruminations that are summarized in Scheme 1. Moore's biosynthesis begins with the tryptophan derivative (**1a**) and terpene (**1b**), which are enzymatically joined *via* chloronium-promoted polyolefin cyclization to provide the tricyclic hapalindole core (*i.e.*, 12-*epi*-hapalindole E (**2**), Scheme 1). At this point, Moore and co-workers proposed that the tricycle can proceed through multiple divergent pathways, the first of which (path A) commences with a cyclization between C(4) of indole and the isopropylidene unit at C(15), leading to the tetracyclic hapalindoles (*i.e.*, 12-*epi*-hapalindole G (**3**)). These natural products can then undergo further oxidation at the indole moiety, leading to the oxindole (*i.e.*, anhydrohapaloxindole A (**4**)), which can be oxidatively cleaved to give the formylkynurenine (*i.e.*, hapalonamide V (**5**)). Alternatively, the tetracyclic hapalindoles can have a *tert*-prenyl moiety attached to C(2) (*i.e.*, ambiguine A (**6**)), which can be engaged in an intramolecular cyclization, leading to the pentacyclic ambiguenes (*i.e.*, ambiguine E (**7**)). These alkaloids can then be further oxidized (*i.e.*, ambiguine D (**8**)) or rearranged (*i.e.*, ambiguine G (**9**)). Furthermore (path B), the tricyclic hapalindoles can undergo cyclization between C(2) and the isopropylidene at C(15), providing the fischerindoles (*i.e.*, 12-*epi*-fischerindole G (**10**)). Finally, Moore and co-workers proposed that the tricyclic hapalindoles can be oxidized to give the putative intermediate **11**, which has not been isolated as a natural product (path C). They further proposed that this intermediate undergoes an acid-catalyzed cyclization to afford welwitindolinone A (**12**), presumably in the pocket of an enzyme. If **12** could be further oxidized, leading to the intermediate epoxide **13**, the [4.3.1]bicyclononanone system could be formed after rearrangement (*i.e.*, "welwitindolinone B isonitrile" (**14a**), which is likely to be a natural product that has yet to be isolated). Moore and co-workers suggested that methylation and oxidation of **14a** could lead to *N*-methylwelwitindolinone C (**15**) and further demonstrated the conversion of **15** into 3-hydroxy-*N*-methylwelwitindolinone C (**17**) and *N*-methylwelwitindolinone D (**19**).

While the Moore biosynthetic hypothesis provides an adequate explanation of the possible relationships between many of the distinct structural classes, a few points remained uncertain, primarily relating to the formation of the welwitindolinones. First, the proposal that **12** arises from the unsaturated intermediate **11** *via* an acid-catalyzed cyclization seems unlikely. The fact that 63 members of this natural product family have been

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Scheme 2. Alternative Biosynthetic Proposal



isolated, while **11** has not been one of them, casts doubt on whether this compound is a plausible intermediate. Given the stability that **11** should demonstrate, at least trace quantities of this compound would be expected in the isolation broths. More importantly, there seems to be little thermodynamic driving force for the conversion of **11** into **12**, due to the generation of a strained spirocyclobutane in this transformation. Second, the mechanistic explanation for the conversion of welwitindolinone A (**12**) into "welwitindolinone B isonitrile" (**14a**) lacks proper literature precedence. Third, the proposal that *N*-methylwelwitindolinone C (**15**) arises from remote oxidation of **14a** could conceivably be explained by an alternate hypothesis. Finally, the isolation literature lacks a biosynthetic hypothesis to account for the genesis of the hapalindolinones.

Given the concerns delineated above, an alternative biosynthetic hypothesis is proposed in Scheme 2. Rather than arising from hypothetical metabolite **11**, welwitindolinone A (**12**) could arise from an oxidative ring contraction of 12-epi-fischerindole I (**21**); the latter could be formed *via* a benzylic oxidation of 12-epi-fischerindole G (**10**), which could in turn arise from the tricyclic hapalindole 12-epi-hapalindole E (**2**). Furthermore, an alternative, albeit untested, mechanistic hypothesis for the conversion of **12** into "welwitindolinone B isonitrile" (**14a**) is put forth that is more in line with literature precedence. Isonitriles are relatively electron-withdrawing, inductively stabilizing negative charges, and have not been invoked as electron-donating entities.<sup>19</sup> It is therefore unlikely that the isonitrile would participate in a fracture of the epoxide *via* electron donation that would lead to a fissure of the cyclobutane ring. Alternatively, based on evidence gleaned in this laboratory,<sup>20</sup>

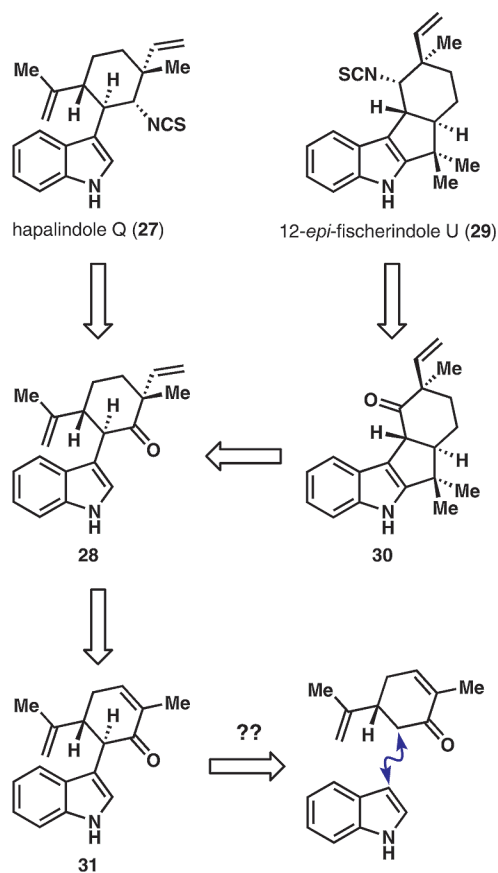
it is more reasonable to invoke electron donation from the N(1) lone pair through the aromatic ring to break the cyclobutane, leading to the  $\alpha$ -isocyanoketone enolate (**22**). This enolate could then attack the highly unstable, extremely electrophilic aza-orthoquinodimethane generated in the reaction, leading to **14a** after tautomerization.<sup>21</sup> Additionally, the remote oxidation of **14a** to *N*-methylwelwitindolinone C (**15**), although perhaps possible with enzymatic intervention, is unlikely. Rather, if **21** were to undergo allylic oxidation, intermediate **23** could be accessed. Upon oxidative ring contraction, similar to that proposed for **12**, the direct product would contain an unstable cyclobutane with vicinal exocyclic olefins. The trisubstituted olefin might then isomerize to form the vinyl chloride (**24**), thus alleviating this additional strain on the cyclobutane. Oxidative ring expansion of **24** and methylation of the indole nitrogen atom could then lead directly to **15**. Finally, we propose that the hapalindolinones could arise from an oxidative coupling event between the C(3) and C(11) carbons of an appropriate tricyclic hapalindole (*i.e.*, 12-epi-hapalindole Q (**25**)), generating the oxindole with the spirocyclopropane moiety (*i.e.*, hapalindolinone B (**26**)).

With a compelling biosynthetic hypothesis in hand, attention could be turned toward designing a synthesis of welwitindolinone A (**12**). However, before such a task was undertaken, it seemed prudent to first consider a synthesis of simpler members of the family, such as hapalindole Q (**27**) and 12-epi-fischerindole U (**29**). It was envisioned that such efforts would reveal hidden clues into the fundamental reactivity of these alkaloids

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**Scheme 3.** Retrosynthetic Analysis of Hapalindole Q (**27**) and 12-*epi*-Fischerindole U (**29**)

and that they would help develop a general route by which to access the core structure. Hapalindole Q (**27**) had been synthesized previously by the Albizati (8 steps, 8.2% overall yield)<sup>10b</sup> and Kerr (12 steps, 1.7% overall yield)<sup>10c,d</sup> groups, but the synthesis of **29** had not been reported prior to our initial communication of this work.<sup>10a</sup> Since a route to **12** was the ultimate goal, a more efficient route to the core (*i.e.*, **27** and **29**) of this alkaloid was required (Scheme 3). Although a cyclization of **27** could certainly be investigated to form **29**, this transformation was instead planned at the ketone stage (**28**), given the acid-sensitivity of the isothiocyanate group. As such, both natural products could be traced back to their respective ketone analogues **28** and **30**, the latter of which should be accessible from the former *via* an acid-catalyzed cyclization. Ketone **28** can be further simplified to the indole/carvone adduct **31** through straightforward functional group transformations. At this stage, a strictly biomimetic synthesis would require a chloronium-promoted polyolefin cyclization to install the terpene moiety; however, a potentially more direct and powerfully simplifying transformation would involve direct formation of the key C(3)–C(10) bond.

### Total Synthesis of Hapalindole Q

Several strategies were investigated for the synthesis of the requisite indole/carvone adduct (**31**) before a method was finally developed to successfully provide the desired material. Initially, it was reasoned that **31** could be accessed from an aldol reaction with subsequent reduction to form the requisite indole moiety (Scheme 4). Indeed, quenching the enolate of carvone with MOM-protected isatin (**32**) provided the desired aldol product

(**33**) in excellent yield. Unfortunately, reduction of this intermediate proved to be problematic. All methods investigated to directly reduce this compound to indole **37** (*e.g.*, LiAlH<sub>4</sub>, BH<sub>3</sub>·THF) were met with failure, so a stepwise solution was sought. Assuming that the C(3) hydroxyl group was preventing reduction of the oxindole, various deoxygenation conditions (*e.g.*, Barton,<sup>22</sup> CS(imid)<sub>2</sub>/hν<sup>23</sup>) were brought to bear for the removal of this alcohol (**38**) prior to indole reduction; however, all attempts were unsuccessful.<sup>24</sup> Reasoning that elimination of the hydroxyl moiety would provide an intermediate upon which further reductions could be performed, various conditions were screened to elicit dehydration (*e.g.*, Martin sulfurane,<sup>25</sup> MsCl/base, TFA/TFAA) before Burgess reagent<sup>26</sup> successfully provided the enone (**36**). However, the yield of this process was irreparably dismal, precluding further chemistry. Since the alcohol could not be reduced, removed, or efficiently eliminated, methods to exchange this moiety for a chlorine atom, which could theoretically be removed with greater facility, were investigated. Employing the conditions developed by Nicolaou,<sup>27</sup> **33** was dissolved in thionyl chloride, providing the cyclic ether **35** (colorless cubes, mp 166–168 °C, see Scheme 4 for X-ray crystallographic analysis) and the unexpectedly stable, semi-protected compound **34**, which led to a dead-end once again.

The resistance of intermediate **33** to reduction necessitated the investigation of alternate means to forge the key bond in **31**. Since removal of the tertiary alcohol had proved to be an intractable problem in the previous route, an approach that circumvented the generation of this group was sought, leading to an oxidative enolate coupling approach to form the desired bond (Scheme 5). The oxidative dimerization of carbonyl compounds has been known for more than 70 years; however, it has yet to become widely utilized by the synthetic community. This reaction has been reviewed,<sup>28</sup> most recently during the full account of the oxidative indole coupling reaction developed in our laboratory,<sup>29</sup> and therefore this information will not be reviewed here. Even though a myriad of literature procedures concerning oxidative enolate couplings was available, many potential drawbacks, could be encountered while pursuing this route. By the inception of this work, only one example of an oxindole oxidative dimerization had been reported.<sup>30a</sup> Additionally, in order to achieve high yields of heterodimerized products

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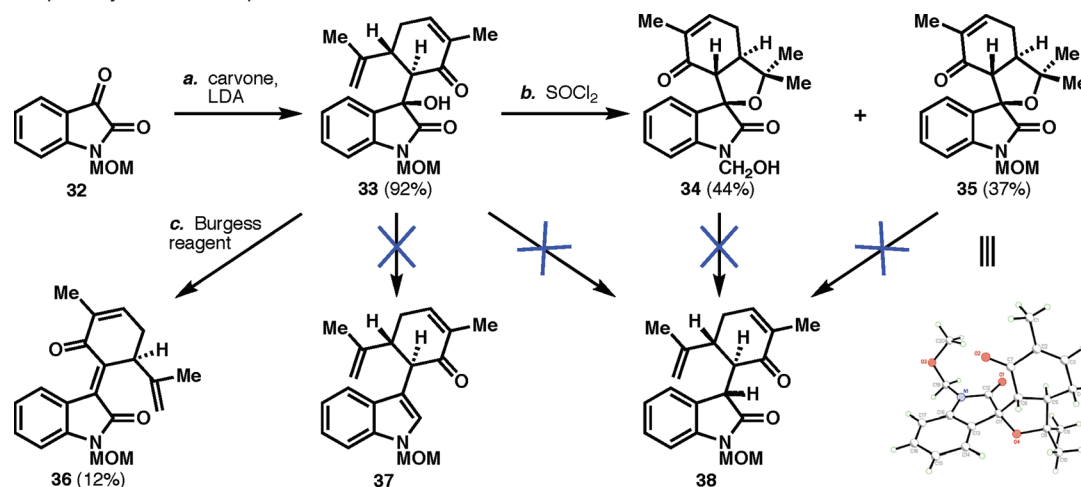
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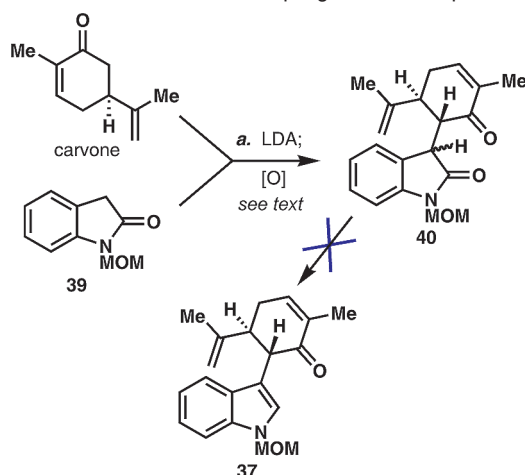
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(30) (a) Fang, C.-L.; Horne, S.; Taylor, N.; Rodrigo, R. *J. Am. Chem. Soc.* **1994**, *116*, 9480–9486. (b) Ito, Y.; Konoike, T.; Saegusa, T. *J. Am. Chem. Soc.* **1975**, *97*, 2912–2914. (c) Ito, Y.; Konoike, T.; Harada, T.; Saegusa, T. *J. Am. Chem. Soc.* **1977**, *99*, 1487–1493.

Scheme 4. Attempted Synthesis of Hapalindole Q via an Aldol Reaction<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a) carvone, LDA (1.0 equiv), THF,  $-78^{\circ}\text{C}$ , 30 min; then **32** (0.83 equiv), 30 min,  $-78$  to  $23^{\circ}\text{C}$ , 92%; (b)  $\text{SOCl}_2$ ,  $23^{\circ}\text{C}$ , 100 min, **34**, 44% and **35**, 37%; (c) Burgess reagent (2.0 equiv), PhH,  $50^{\circ}\text{C}$ , 3 h, 12%. LDA, lithium diisopropylamide; THF, tetrahydrofuran; PhH, benzene.

Scheme 5. Oxindole Oxidative Coupling Route to Hapalindole Q<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a) carvone (1.0 equiv), **39** (1.0 equiv), LDA (2.1 equiv), THF,  $-78^{\circ}\text{C}$ , 30 min; then oxidant (2.0 equiv),  $23^{\circ}\text{C}$ , 20 min. LDA, lithium diisopropylamide; THF, tetrahydrofuran; [O], oxidation.

in an oxidative enolate coupling, 3 equiv or more of one coupling partner was usually required.<sup>30b,c</sup> Finally, few investigations into the factors that govern the heterodimerization event had been performed; thus, it was unknown whether any of the desired heterocoupled product would be obtained. Despite such potential drawbacks, an examination of the oxidative oxindole coupling was undertaken, which would at least provide further insight into the elements that govern such transformations.

As a starting point for the examination of this heterocoupling reaction, treatment of carvone and MOM-protected oxindole (**39**)<sup>31</sup> with  $\text{FeCl}_3$  in DMF provided the desired product (**40**) in *ca.* 15% yield. Given the success of this coupling, an optimization of this particular reaction was undertaken, centering primarily on oxidant selection (Table 1). Hypervalent iodine and cupric-based oxidants were less efficient at promoting the coupling than the ferric-based systems, so the focus of further studies was narrowed. Success was finally realized when acetylacetonate-type ligands were utilized on the iron center.

Table 1. Optimization of the Oxidative Oxindole Coupling<sup>a</sup>

oxidant	yield (%)
$\text{PhI}(\text{OAc})_2$	10
$\text{Cu}(\text{2-ethylhexanoate})_2$	15
$\text{FeCl}_3$	~15
$\text{Fe}(\text{PhCOCHCOCH}_3)_3$	30
$\text{Fe}(\text{CH}_3\text{COCHCOCF}_3)_3$	40
$\text{Fe}(\text{C}_{10}\text{H}_7\text{COCHCOCF}_3)_3$	40
$\text{Fe}(\text{CH}_3\text{COCHCOCH}_3)_3$	45
$\text{Fe}(t\text{-BuCOCHCOCH}_3)_3$	83

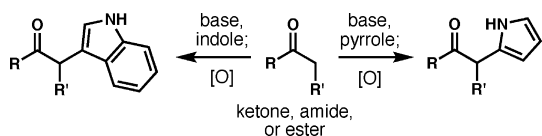
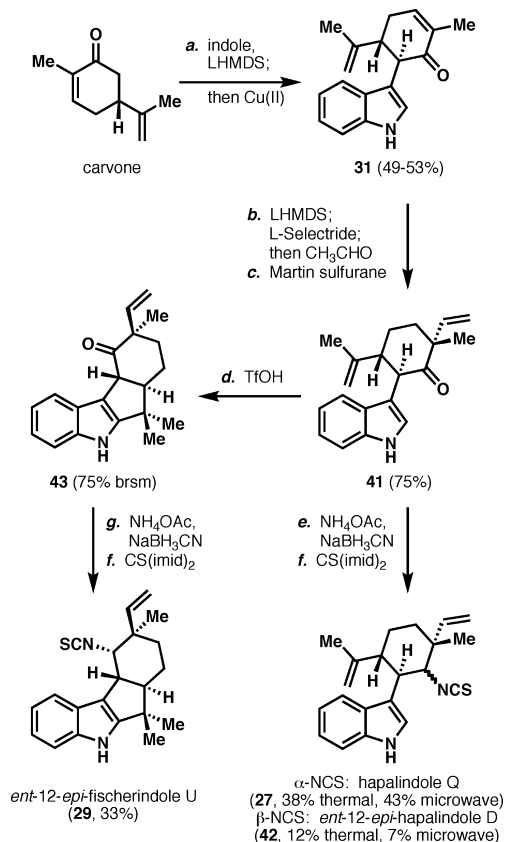
<sup>a</sup> LDA, lithium diisopropylamide; [O], oxidation.

Specifically,  $\text{Fe}(t\text{-BuCOCHCOCH}_3)_3$  was exceptionally efficient, providing an 83% isolated yield of the desired product (as a 1:1 mixture of diastereomers at C(3)), utilizing equimolar quantities of both coupling partners! These results have led to the hypothesis that careful tuning of the oxidant's oxidation potential to more closely match one of the coupling partners could lead to selective heterodimerizations.<sup>32</sup> With high-yielding access to intermediate **40**, efforts were turned toward the reduction of oxindole **40** to indole **37** (Scheme 5). Once again, the reduction proved recalcitrant, necessitating yet another reevaluation of the synthetic strategy.

As with the aldol route, an alternative to the oxindole coupling was sought: one that would avoid a problematic reduction step. As such, the most straightforward method by which to circumvent this difficulty would be to directly attach the indole to the carvone moiety. Unfortunately, no such method had been reported, requiring the invention of chemistry to fill this gap. Inspired by the oxidative coupling literature, in conjunction with Barton's classic synthesis of usnic acid,<sup>33</sup> the oxidative indole coupling was conceived. In this reaction,<sup>10a,29</sup> an indole can be

(31) Wang, J.-J.; Hu, W.-P. *J. Org. Chem.* **1999**, *64*, 5725–5727.

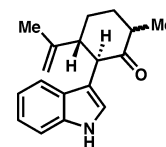
## Scheme 6. Oxidative Indole and Pyrrole Couplings

Scheme 7. Total Syntheses of Hapalindole Q (27), *ent*-12-*epi*-Hapalindole D (42), and *ent*-12-*epi*-Fischerindole U Isothiocyanate (29)<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a) indole (2.0 equiv), carvone (1.0 equiv), LHMDS (3.3 equiv), THF,  $-78^{\circ}\text{C}$ , 30 min; then copper(II) 2-ethylhexanoate (1.5 equiv),  $-78$  to  $23^{\circ}\text{C}$ , 15 min, 49–53%; (b) LHMDS (1.5 equiv), THF,  $-78^{\circ}\text{C}$ , 20 min; then L-Selectride (1.05 equiv), 1 h, then  $\text{CH}_3\text{CHO}$  (6.0 equiv),  $-78$  to  $23^{\circ}\text{C}$ , 2 h; (c) Martin sulfurane (1.1 equiv),  $\text{CHCl}_3$ , 10 min, 75% (2 steps); (d) TMSOTf (3 equiv), MeOH (1.1 equiv), DCM,  $0^{\circ}\text{C}$ , 1 h, 31% isolated, 75% brsm; (e)  $\text{NaCNBH}_3$  (10 equiv),  $\text{NH}_4\text{OAc}$  (40 equiv), MeOH, THF, microwave,  $150^{\circ}\text{C}$ , 2 min, 61% combined; (f)  $\text{CS}(\text{imid})_2$  (1.1 equiv), DCM,  $0$  to  $23^{\circ}\text{C}$ , 3 h; (g)  $\text{NaCNBH}_3$  (10 equiv),  $\text{NH}_4\text{OAc}$  (40 equiv), MeOH, THF,  $23^{\circ}\text{C}$ , 7 d, 55%. LHMDS, lithium hexamethyldisilazide; THF, tetrahydrofuran; TMSOTf, trimethylsilyl trifluoromethanesulfonate; DCM, dichloromethane.

directly attached to a variety of carbonyl compounds in good yields (Scheme 6). The method was found to generate the desired indole/carvone adduct (**31**, colorless cubes, mp  $129$ – $130^{\circ}\text{C}$ , structure verified by X-ray crystallographic analysis) in good yield and in one synthetic operation from commodity starting materials (Scheme 7); this has also been extended to the coupling of unfunctionalized pyrroles.<sup>34</sup>

## Chart 2. Reductive Alkylation Byproduct



With gram quantities of **31** in hand, attention was turned to the completion of the total synthesis of **27** and **29**, which first required reductive alkylation of the C(12)–C(13) olefin.<sup>35</sup> Treatment of **31** with 2 equiv of L-Selectride was expected to first deprotonate N(1)–H and then perform a conjugate reduction to generate the enolate, which could subsequently be trapped with acetaldehyde. Surprisingly, 1,4-hydride addition preceded N(1)–H deprotonation, thus generating the enolate, which in turn deprotonated N(1), forming the corresponding reduced enone (see Chart 2). Rather than protecting the indole nitrogen atom, N(1)–H was first deprotonated with LHMDS, followed by addition of L-Selectride, and the resulting enolate was trapped with acetaldehyde. This intermediate alcohol was immediately dehydrated with Martin sulfurane to give vinylated compound **41**, which intersected the Albizati synthesis.<sup>10b</sup> Reductive amination using Albizati's conditions ( $\text{NH}_4\text{OAc}$ ,  $\text{NaCNBH}_3$ , room temperature, 7 days) provided the desired amine in 61% yield as a 3:1 mixture of diastereomers. Alternatively, microwave irradiation ( $150^{\circ}\text{C}$ , 2 min) provided an identical yield of the product, with an increased diastereomeric ratio of 6:1. To the best of our knowledge, this is a unique example of an increase in diastereoselectivity as a consequence of microwave irradiation. Treatment of the resulting amine with  $\text{CS}(\text{imid})_2$  installed the isothiocyanate, thus completing the total synthesis of **27** and the first total synthesis of *ent*-12-*epi*-hapalindole D (**42**). Tricyclic ketone **41** could also undergo a biomimetic<sup>7</sup> acid-catalyzed cyclization to provide the tetracyclic ketone **43** upon exposure to *in situ*-generated triflic acid.<sup>36</sup> Reductive amination and formation of the isothiocyanate completed the first total synthesis of **29**, which also allowed the determination of the absolute stereochemistry of the fischerindole-type natural products (*i.e.*, opposite to that depicted in Scheme 7). The syntheses of **27**, **42**, and **29** proceeded enantiospecifically in 15%, 4.8%, and 9.8% overall yields, respectively, without resorting to protecting groups or superfluous redox manipulations.<sup>10a</sup>

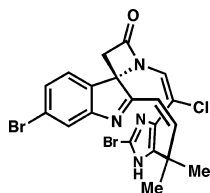
## Retrosynthetic Analysis of Welwitindolinone A

Having demonstrated that the direct indole coupling could enable rapid access to several of the simpler members of the hapalindole family of natural products, attention could then be turned to the more complex members. Welwitindolinone A (**12**) was an attractive target for total synthesis due to its strikingly unique molecular architecture. It contains three all-carbon quaternary centers, two of which are chiral, and an asymmetrically disposed neopentyl chlorine atom, all incorporated into a highly strained spirocyclobutane-containing oxindole. As already discussed (*vide supra*), the revised biosynthetic hypothesis proposes that 12-*epi*-fischerindole I (**21**) is converted into **12** *via* oxidative ring contraction.<sup>37</sup> Such ring contractions are primarily utilized for the preparation of spirocyclic five-membered rings from annulated six-membered rings and often

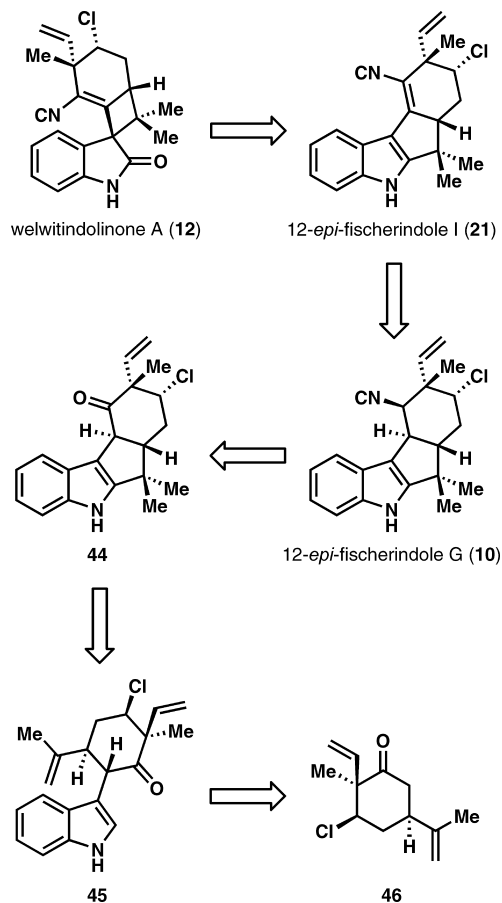
(32) (a) Baran, P. S.; DeMartino, M. P. *Angew. Chem., Int. Ed.* **2006**, *45*, 7083–7086. (b) DeMartino, M. P.; Chen, K.; Baran, P. S. *J. Am. Chem. Soc.* **2008**, *130*, 11546–11560.  
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(35) Fortunato, J. M.; Ganem, B. *J. Org. Chem.* **1976**, *41*, 2194–2200.  
(36) Prolonged exposure to triflic acid led to product decomposition; therefore, the reaction was quenched early and the starting material was recycled.

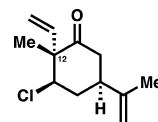


Chart 3. Strained  $\beta$ -Lactam in Chartelline C

## Scheme 8. Retrosynthetic Analysis of Welwitindolinone A (12)



require relatively harsh reaction conditions to proceed.<sup>38</sup> Only one example has been reported for the direct preparation of a spirocyclic four-membered ring from an annulated five-membered ring;<sup>39</sup> however, a related conversion has been observed in this laboratory to generate a strained  $\beta$ -lactam (see Chart 3).<sup>20</sup> Furthermore, it is conceivable that **21** could arise from 12-*epi*-fischerindole G (**10**) via benzylic oxidation (Scheme 8). Through straightforward functional group manipulations, **10** could be derived from the tetracyclic ketone **44**, which in turn could arise from an acid-catalyzed cyclization of tricyclic ketone **45**. Exploiting the direct coupling methodology developed for the synthesis of **27**, **45** could be obtained from the coupling of indole with **46**. Chloroketone **46** was a new chemical entity; however, a very similar analogue had been prepared by Fukuyama *en route* to hapalindole G.<sup>7</sup> Fukuyama's chloroketone

Chart 4. Fukuyama's Chloroketone: Diastereomer of **46** at C(12)

(see Chart 4) is diastereomeric at C(12), and the chemistry utilized for its preparation was unfortunately not amenable for the preparation of **46**.

Total Synthesis of *ent*-Fischerindole G

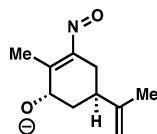
Historically, installation of chlorine atoms adjacent to quaternary centers has been met with difficulty, and it was anticipated that **46** would not be an exception.<sup>40</sup> Forays into the preparation of this compound focused on direct means for concurrent installation of the chlorine atom and the quaternary stereocenter. As such, attempts were made to apply a modified Baylis–Hillman-type reaction<sup>41</sup> in which  $\beta$ -chloroketones are generated from the corresponding enones (Scheme 9), with concomitant incorporation of an aldehyde. However, this reaction was never utilized to install a quaternary center at the  $\alpha$ -position prior to this work. Unfortunately, none of the desired product (**52**) was obtained using these conditions. Given the limited prospects for direct installation of the hallmark chlorine atom, attention was instead focused on a variety of chlorine equivalents. Carboxylic acids can be readily converted into chlorines *via* a Hunsdiecker reaction, so several acyl anion equivalents (*e.g.*, 1,3-dithiane derivatives, (MeO)<sub>3</sub>CH, (PhS)<sub>3</sub>CH) were examined in 1,4-additions to carvone, but none of them provided the desired product. Encouragingly, successful 1,4-addition was observed with phenylcuprate, and the incipient enolate was trapped with acetaldehyde to give **47**. However, attempts to oxidatively degrade the aromatic ring<sup>42</sup> led to significant overoxidation. A thiophenyl group was next targeted as a chlorine equivalent, due to the propensity of such moieties to undergo  $\alpha$ -chlorination or Pummerer rearrangements. Thiophenol participated in an efficient vicinal difunctionalization of carvone with aluminum catalysis<sup>43</sup> to provide, after dehydration with Martin sulfurane, thioether **49**. Attempts to directly chlorinate **49** were unsuccessful using a variety of reagents (*e.g.*, NCS, Raney Ni/CCl<sub>4</sub>, SO<sub>2</sub>Cl<sub>2</sub>, CCl<sub>4</sub>/*h* $\nu$ , Cl<sub>3</sub>CCOCCl<sub>3</sub>). Chlorination with concomitant removal of the thiol under numerous conditions (*e.g.*, PIFA/LiCl, LiNaphthalenide/TsCl, TiCl<sub>4</sub>/CCl<sub>4</sub>, MeI/NaCl<sup>44</sup>) was also investigated, but to no avail. Routes in which the ketone was first converted into the chloride, followed by sulfide oxidation, were also considered. Direct reductive chlorination of the ketone,<sup>45</sup> instead of providing the desired product, led to the unexpected bicycle **50**. Attempts to generate the vinyl chloride *via* POCl<sub>3</sub>, which could potentially be carefully hydrogenated to the alkyl chloride, were also fruitless. Unable to perform a direct reductive chlorination of the carbonyl, **49** was first reduced to the neopentyl alcohol (**51**) and then unsuccessfully subjected to a variety of chlorination

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 (45) Onishi, Y.; Ogawa, D.; Yasuda, M.; Baba, A. *J. Am. Chem. Soc.* **2002**, *124*, 13690–13691.



## Chart 5. Anticipated Vinylnitroso Intermediate



and co-workers.<sup>53</sup> Reaction of the oxime with 2 equiv of a cuprate reagent would first deprotonate the oxime, which could then open the epoxide and lead to the corresponding vinylnitroso compound (see Chart 5). The second equivalent of cuprate reagent could then participate in a 1,4-addition into the vinylnitroso group, thereby installing the quaternary center. Unfortunately, divinylcuprate was unsuccessful at accomplishing this transformation.

After several other failed attempts to generate the desired chloroketone **46**, a successful route was finally developed, inspired by a reaction developed in the Wender laboratory.<sup>54</sup> In Wender's study,  $\alpha,\beta$ -epoxyketones were first treated with strong base to form the corresponding enolate. Nucleophilic addition of an organometallic reagent (usually Grignard reagents) to this epoxyenolate at the  $\alpha$ -carbon (as opposed to the usual  $\beta$ -attack) formed the corresponding  $\alpha$ -alkyl- $\beta$ -hydroxyketone, and subsequent elimination of the hydroxyl group furnished the  $\alpha$ -substituted enone. However, a quaternary carbon installation during the course of this reaction was unprecedented. Despite this potential limitation, the reaction was attempted on carvone oxide to provide **59** in about 30% yield (Scheme 10). Extensive optimization efforts (solvent, nucleophile, base, additives, temperature, addition rates) did not result in significant improvement in the overall efficiency of the reaction. The low yield is likely due to the sterically hindered nature at the  $\alpha$ -position of the  $\alpha,\beta$ -epoxyketone, causing  $S_N2'$  attack (**58**, red arrow) to be favored over the desired  $S_N2$  attack (**58**, blue arrow).<sup>55</sup> Nevertheless, with alcohol **59** readily available, the chlorination was accomplished (NCS/ $PPh_3$ ) in acceptable yield to provide the key chloroketone (**46**).<sup>56</sup> Despite the modest overall yield of this two-step sequence, it can be used to rapidly prepare multigram quantities of **46** and was therefore deemed as an acceptable solution to this problem, especially given the difficulties in preparing **46** *via* other routes (*vide supra*). It is noteworthy that **59** was the only neopentyl alcohol that could be successfully chlorinated during the course of these studies. It is also interesting that varying amounts of the diastereomer (at C(13)) are generated, although these chlorination conditions often proceed with complete inversion of stereochemistry. Perhaps this alcohol is more accessible to electrophiles due to a slight bond-lengthening effect (*i.e.*, *retro*-aldol ability), thus mitigating the neopentyl hindrance and allowing epimerization to a minor extent.

With **46** in hand, the stage was finally set to invoke the direct indole coupling reaction, which proceeded smoothly to provide the coupled product (**45**) in 62% yield as a single diastereomer (verified by X-ray crystallographic analysis). Several aspects

of this particular coupling are noteworthy. First, it is remarkable that chloride elimination is not observed during the course of this coupling. Second, it was discovered that, as the reaction concentration was increased, the yield improved. Third, any C(13) diastereomer of **46** present in the reaction mixture does not participate in the coupling reaction and rather suffers elimination, presumably due to the axial disposition of the chlorine atom. Finally, this effective method for direct C–C bond formation enables all the necessary carbon atoms of these complex natural products to be secured in only three steps and was routinely carried out on multigram scale. Functional group manipulations are all that remained to complete the synthesis of **12**.

Conditions (*i.e.*, TfOH) previously applied to the conversion of **41** to **43** unfortunately provided significant quantities of several byproducts in the attempted conversion of **45** to **44**,<sup>57</sup> including **65**, **66**, and **67** (Scheme 11). In this particular cyclization reaction, two modes of activation are feasible at the *gem*-disubstituted olefin. This olefin can be protonated to give the tertiary carbocation, which is intercepted by the indole ring to give intermediate **61**, which in turn leads to the desired product (**44**). Alternatively, protonation to give the primary carbocation, which is also intercepted by the indole ring, leads to intermediate **62**. A [1,5]-sigmatropic shift generates **63**, which can then rearrange to carbocation **64**. The three possible modes of elimination to quench this carbocation provide the three observed products (**65**, **66**, and **67**; **67** as clear cubes, mp 128–129 °C, see Scheme 11 for X-ray crystallographic analysis). In order to circumvent the formation of such undesired byproducts, a variety of Lewis and Brønsted acids were screened (TFA, HCl,  $MeOSO_2H$ ,  $H_2SO_4$ , heat, TsOH,  $BF_3 \cdot Et_2O$ , AcOH,  $Tf_2NH$ ,  $Dy(OTf)_3$ , PPTS,  $AlCl_3$ ,  $FeCl_3$ , silicotungstic acid,  $PtCl_2$ , zeolite NaY,  $RuCl_3/AgOTf$ ,  $Cu(OTf)_2$ ,  $Pd(OAc)_2$ ,  $Co(acac)_2/PhSiH_3$ ), many of which did not catalyze the cyclization and none of which provided either any improvement in the yield or reduction in the quantity of byproducts. Attempted cyclizations of the amine or alcohol derivatives of **45** were equally unsuccessful. Finally, it was found that Montmorillonite K-10 acidic clay, with microwave irradiation, provided the desired product (**44**) without formation of the undesired byproducts, although recycling of unreacted starting material was required. It was assumed, based on the reactivity observed for **43**, that reductive amination of the ketone **44** would provide amine **60** directly. However, reductive amination under the previously employed conditions (see Scheme 7)<sup>58</sup> provided amine **70** (see Scheme 13) as a single diastereomer, which was epimeric at C(11) (as was required for **10**). This unexpected complete inversion in diastereoselectivity seems to be solely due to the presence of the C(13) chlorine atom. A more circuitous route was therefore required to access the desired amine (**60**), necessitating reduction to the alcohol, mesylation, azidation, and reduction (Scheme 10), similar to the sequence developed by Fukuyama and co-workers for hapalindole G.<sup>7</sup> Formylation<sup>59</sup> of the amine (**60**) followed by dehydration with Burgess reagent<sup>60</sup> provided *ent*-12-*epi*-fischerindole G (**10**),<sup>16</sup> which was spectroscopically identical to the natural material with the exception of optical rotation. Thus, in principle, the naturally

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(54) Wender, P. A.; Erhardt, J. M.; Letendre, L. J. *J. Am. Chem. Soc.* **1981**, *103*, 2114–2116.

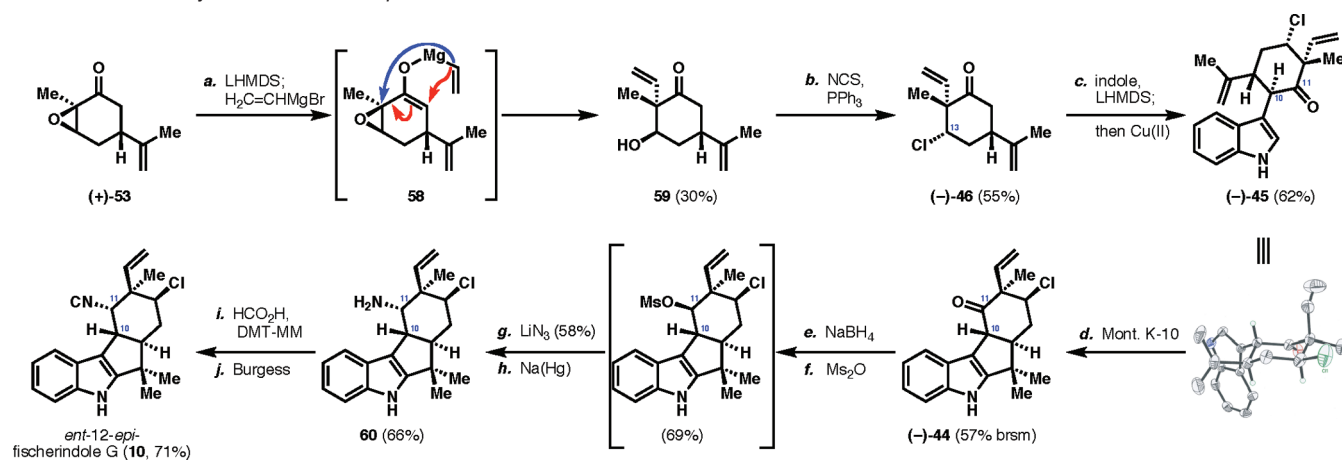
(55) This alternate reaction pathway was also noted in Wender's studies.

(56) Alternative chlorination conditions were attempted ( $SOCl_2$ /base,  $PPh_3/CCl_4$ ,  $PPh_3/ZnCl_2/DEAD$ ,  $MsCl/Pyrr.$ ,  $PPh_3/Cl_2$ , DMAP/CS(imid)<sub>2</sub>,  $(COCl)_2$ , TCT,  $TMSCl/BiCl_3$ ) but were unsuccessful at providing the desired product, and *retro*-aldol product was observed on at least one occasion.

(57) In fact, the reactivity of **47** and downstream intermediates differed significantly from that observed with the 12-*epi*-fischerindole U series.

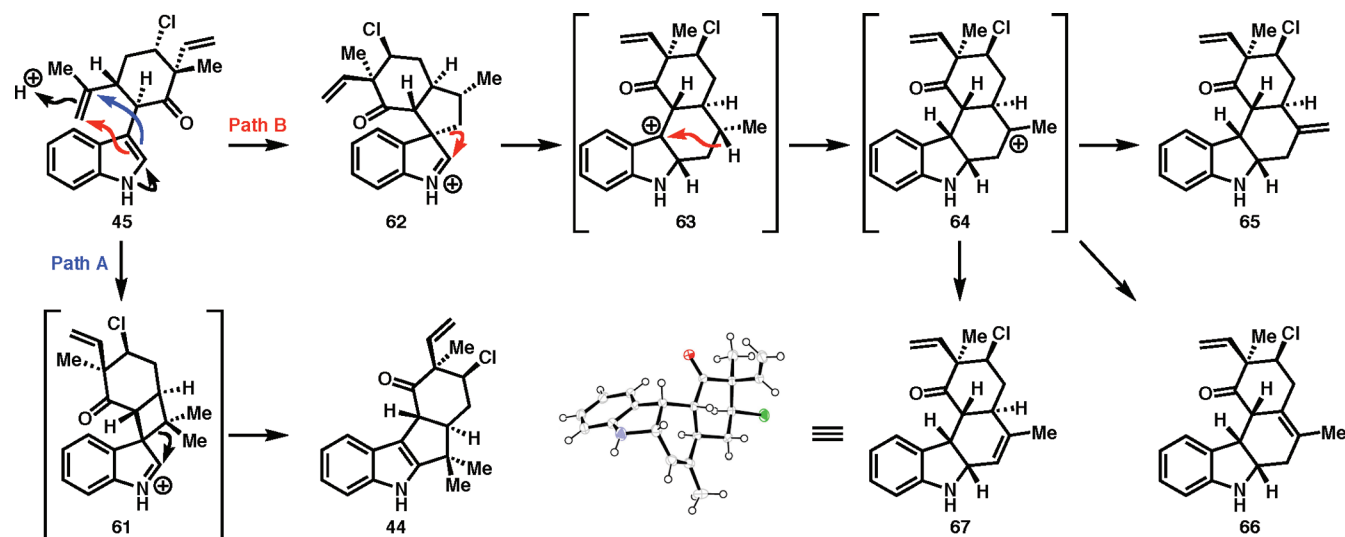
(58) Attempts to utilize microwave irradiation to promote this reaction led to dechlorination of the compound and were therefore not amenable for this series of chlorinated natural products.

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Scheme 10. Total Synthesis of *ent*-12-*epi*-Fischerindole G<sup>a</sup>

occurring enantiomer of **10** could be prepared from (*S*)-carvone oxide (*vide infra*).<sup>61</sup>

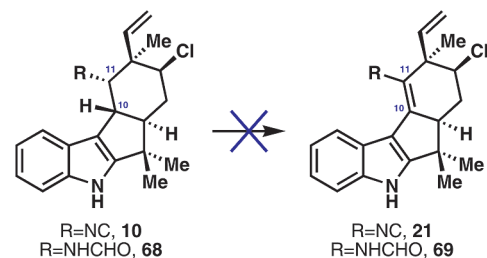
Scheme 11. Formation of the Acid-Catalyzed Cyclization Byproducts



occurring enantiomer of **10** could be prepared from (*S*)-carvone oxide (*vide infra*).<sup>61</sup>

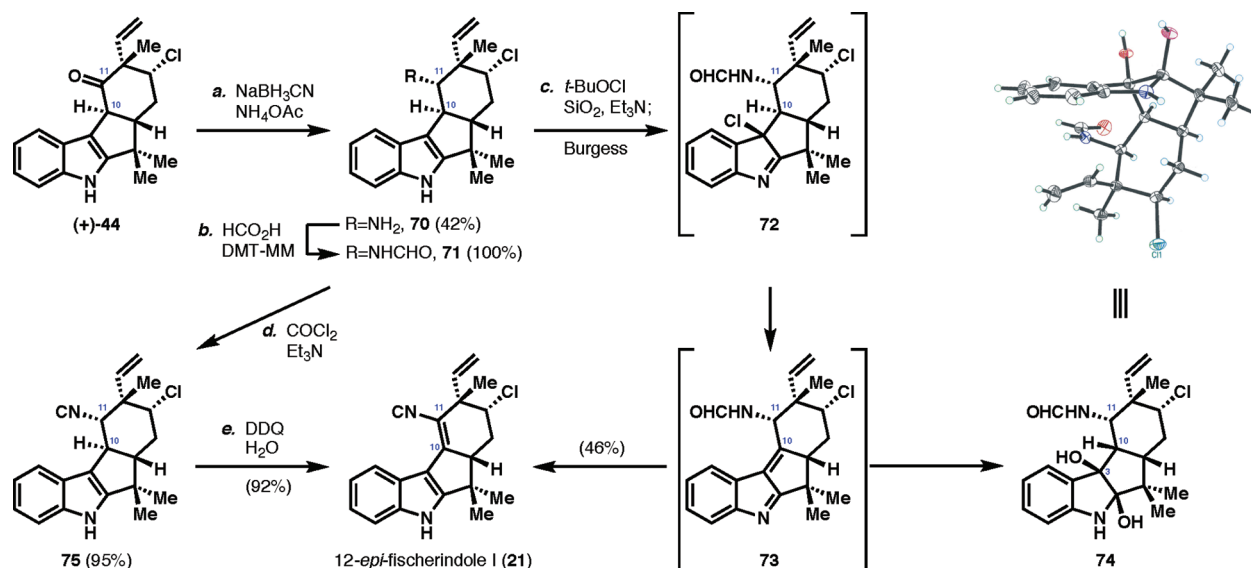
### Total Synthesis of Fischerindole I

With reasonable quantities of **10** available, the oxidation to form 12-*epi*-fischerindole I (**21**) could be investigated. Treatment of **10** with a variety of oxidants (*e.g.*, DDQ,  $\text{MnO}_2$ , *p*-chloranil, *t*-BuOCl) failed to produce any **21** (Scheme 12). Reasoning that the isonitrile could preferentially react with these oxidants, the formamide derivative (**68**) was subjected to various oxidants,<sup>62</sup> but this also failed to form any of the desired product. Putative formation of stable 3-chloro- or 3-hydroxyindolenines was the only reactivity observed with either **10** or **68**, but these could not be utilized to install the required element of unsaturation. Turning to the amine diastereomer **70** (Scheme 13),<sup>63</sup> which

Scheme 12. Attempted Conversion of *ent*-12-*epi*-Fischerindole G (**10**) into *ent*-12-*epi*-Fischerindole I (**21**)

was ineffective for the preparation of **10**, it was hoped that this intermediate would allow the desired oxidation to proceed through subtle stereoelectronic differences. Amine **70** was formylated in quantitative yield to give **71**, which was then subjected to various oxidation conditions.<sup>64</sup> Remarkably, **21** was produced in 46% overall yield when formamide **71** was treated

(60) Creedon, S. M.; Crowley, H. K.; McCarthy, D. G. *J. Chem. Soc., Perkin Trans. 1* **1998**, 1015–1017.

Scheme 13. Original and Improved Total Synthesis of 12-*epi*-Fischerindole I (**21**)<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a)  $\text{NH}_4\text{OAc}$  (40 equiv),  $\text{NaCNBH}_3$  (7.5 equiv), 3 Å molecular sieves, MeOH, THF, sonication, 18 h, 42%; (b)  $\text{HCO}_2\text{H}$  (2.0 equiv), CDMT (2.2 equiv), DMAP (0.1 equiv), NMM (2.2 equiv), DCM, 23 °C, 30 min, 100%; (c) THF, TEA (1.0 equiv), *t*-BuOCl (1.5 equiv), 0 °C, 10 min, then  $\text{SiO}_2/\text{TEA}$  (PTLC); then  $\text{CDCl}_3$ , then Burgess reagent (2.0 equiv), PhH, 23 °C, 30 min, 46% overall; (d) TEA (17.5 equiv), DCM,  $\text{COCl}_2$  (2.0 equiv), 0 °C, 10 min, 95%; (e) DDQ (2.5 equiv),  $\text{H}_2\text{O}$ , THF, 0 °C, 30 min, 92%. THF, tetrahydrofuran; CDMT, 2-chloro-4,6-dimethoxy-1,3,5-triazine; DMAP, 4-(dimethylamino)pyridine; NMM, *N*-methylmorpholine; DCM, dichloromethane; DMT-MM, 4-(4,6-dimethoxy-1,3,5-triazin-2-yl)-4-methylmorpholinium chloride; TEA, triethylamine; PTLC, preparative thin-layer chromatography; PhH, benzene; DDQ, 2,3-dichloro-5,6-dicyanobenzoquinone.

first with *t*-BuOCl and triethylamine and subsequently with Burgess reagent. Although the intermediates in this reaction are unstable and difficult to purify, it is reasonable to assume that the reaction proceeds *via* the intermediates delineated in Scheme 13. Initial chlorination of the indole leads to chloroindolenine **72**, which undergoes elimination to generate methylene indolenine **73**. This intermediate can tautomerize to 12-*epi*-fischerindole I formamide (**69**), which is unstable and immediately dehydrates upon exposure to Burgess reagent to provide **21**.<sup>16</sup> Methylene indolenine **73** can alternatively be hydrated (presumably on silica gel) at the imine carbon. Reprotonation of the olefin from the  $\beta$ -face and attack by water at C(3), *via* the intermediacy of an azaorthoquinodimethane intermediate, provides the major side product (**74**, confirmed by X-ray crystallographic analysis). It is to be noted that the presence of the epimerized C(10) center provides evidence for the existence of **73 en route** to **21**.

Although this sequence provided the first synthetic sample of **21**, it was plagued with numerous problems, specifically low overall yields, difficult and unscalable synthetic procedures, the formation of byproducts, and the intermediacy of several unstable intermediates. It was reasoned that the low overall efficiency of this route likely stemmed from two controlling factors. First, *t*-BuOCl is not commonly employed as an oxidant for benzylic oxidations. Perhaps an oxidant more chemoselective for such transformations would provide a higher yield of the desired product. Second, the penultimate intermediate (**69**, Figure 1) in this sequence is an extremely unstable compound

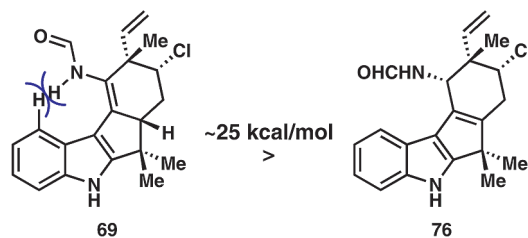


Figure 1. Calculated stability of the 12-*epi*-fischerindole I penultimate intermediate.

due to a severe steric interaction between the formamide N–H and the C(4)–H of the indole ring, leading to an approximate 25 kcal/mol calculated destabilization compared to its olefin-translocated isomer (**76**).<sup>65</sup> Therefore, an alternate sequence was sought to accomplish this transformation while bypassing these difficulties. It was subsequently discovered that initial dehydration of formamide **71** (Scheme 13) provided isonitrile **75**, which is epimeric at C(11) to 12-*epi*-fischerindole G (**10**), in 95% yield. Treatment of this compound with DDQ,<sup>66</sup> in contrast to the DDQ reaction for **10**, provided 12-*epi*-fischerindole I (**21**) in 92% yield, allowing access to large quantities of this natural product in short order.<sup>14</sup> Based on these results, although it is quite possible that **10** could be enzymatically promoted to **21**, it is equally possible that **75** (not yet isolated as a natural product), rather than **10**, could be the actual biosynthetic precursor to **21**.

### Total Synthesis of Welwitindolinone A

With 12-*epi*-fischerindole I (**21**) in hand, it was finally possible to investigate the conversion of this natural product into welwitindolinone A (**12**). Although the proposed conversion of **21** to **12** might appear intuitive, practical difficulties were

(61) Until this point, for initial studies into the synthesis of these natural products, (*R*)-carvone oxide was used because of its significantly lower cost.

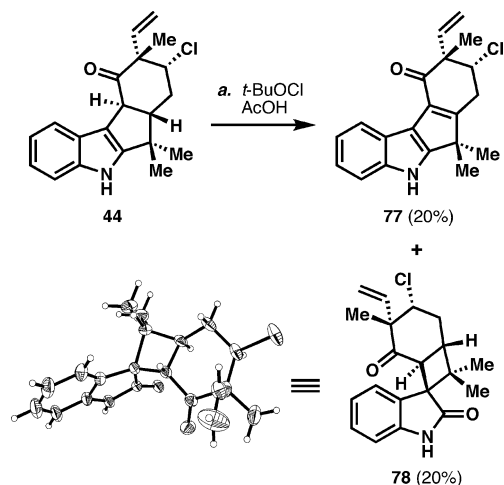
(62) The same oxidants were utilized as for **10**, as well as DMDO and DMP.

(63) From this point on, (*S*)-carvone oxide was utilized to allow access to the correct enantiomer of the natural product.

(64) It was assumed that the isonitrile might react disastrously with various oxidants; therefore, the formamide was utilized in these trials.

(65) As determined by AM1 calculations.

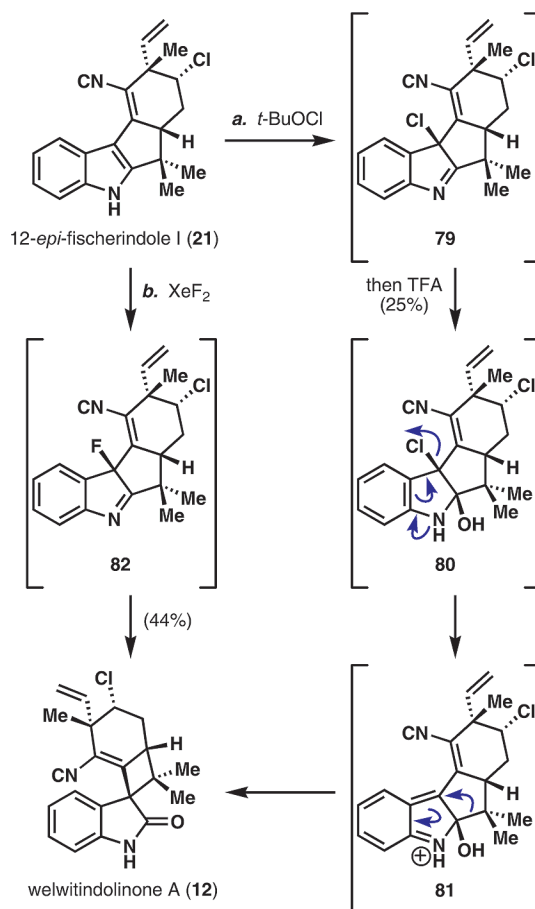
(66) Oikawa, Y.; Yonemitsu, O. *J. Org. Chem.* **1977**, *42*, 1213–1216.

**Scheme 14.** Model System for the Total Synthesis of Welwitindolinone A (**12**)<sup>a</sup>

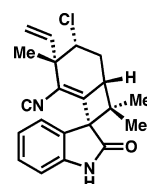
<sup>a</sup> Reagents and conditions: (a) THF, TEA (1.0 equiv), *t*-BuOCl (1.7 equiv), 0 °C, 10 min; then 40:20:1 MeOH:H<sub>2</sub>O:AcOH, 5 min. THF, tetrahydrofuran; TEA, triethylamine.

expected due to the sensitivity of both alkaloids to acidic media and the sheer ring strain of the resulting product. Such concerns were intensified by the knowledge that oxidative ring contractions to generate five- and six-membered rings typically require elevated temperatures, and hence, forming a strained four-membered ring should be even more difficult.<sup>37</sup> Before using valuable material to probe this transformation, a model system was sought on which to test this ring contraction. To our delight, treatment of cyclized ketone **44** with *t*-BuOCl and then dilute AcOH furnished two major products: the oxidized compound **77** and ring-contracted compound **78** in ~20% unoptimized yield each (Scheme 14). Given the aforementioned considerations, it is remarkable that this reaction occurs at low temperature and within minutes. Although ketone **78** represents a potentially viable intermediate to complete the synthesis of **12**,<sup>67</sup> attention was returned to **21** in the hope that it could be directly converted into **12**, thus lending credence to the proposed biosynthetic hypothesis (*vide supra*).

Extensive experimentation led to conditions by which **21** could be converted directly into **12**. 12-*epi*-Fischerindole I (**21**) was exposed to *t*-BuOCl and triethylamine in THF at -30 °C for 1 min, and the solvent was rapidly removed (Scheme 15). The crude residue was then dissolved in a THF:H<sub>2</sub>O:TFA mixture (95:4:1) and warmed to 0 °C. Strict adherence to this protocol resulted in the first total synthesis of **12** in 25% yield,<sup>16</sup> along with minor amounts of its C(3) epimer (see Chart 6). This transformation most likely proceeds through initial chlorination of the indole to provide **79**, followed by attack of water to give **80**. Elimination of the chloride to generate azaorthoquinodimethane **81**, followed by [1,5]-sigmatropic rearrangement, would then install the spirocyclobutane of **12**. Although **12** was accessible *via* this procedure, there were several problems with this route, most notably the low yield and the exclusive formation of the C(3) diastereomer upon scale-up. Seeking to circumvent these obstacles, it was determined that the chief difficulty arose from the isonitrile moiety, which has been shown to be unstable to electrophilic chlorinating reagents in this

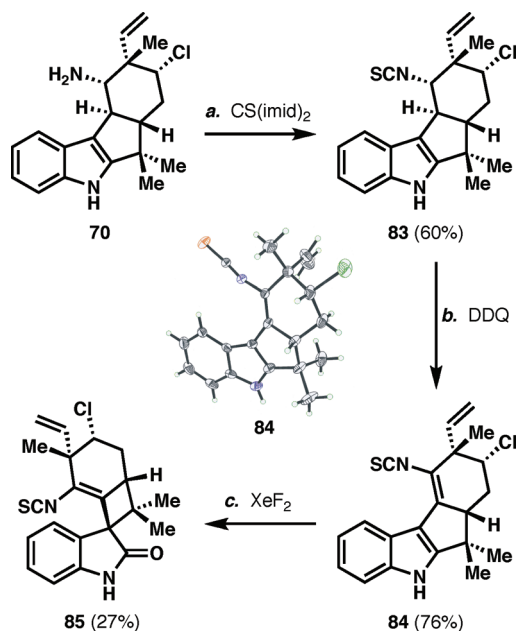
**Scheme 15.** Original and Improved Total Syntheses of Welwitindolinone A (**12**)<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a) THF, TEA (1.0 equiv), *t*-BuOCl (1.5 equiv), -30 °C, 1 min; then 95:4:1 THF:H<sub>2</sub>O:TFA, -30 to 0 °C, 5 min, 25%; (b) XeF<sub>2</sub> (1.0 equiv), H<sub>2</sub>O, MeCN, 23 °C, 5 min, 44%. THF, tetrahydrofuran; TEA, triethylamine; TFA, trifluoroacetic acid; MeCN, acetonitrile.

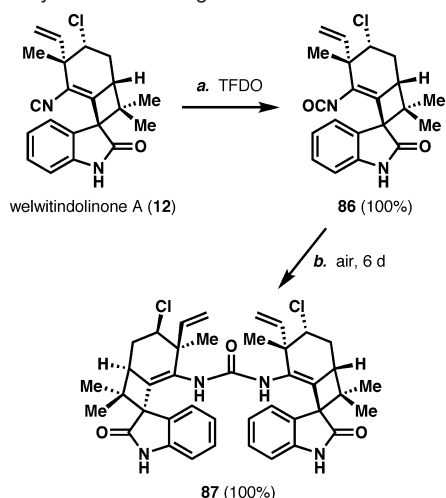
**Chart 6.** 3-*epi*-Welwitindolinone A

laboratory.<sup>14</sup> It was reasoned that a hitherto-unknown fluoro-hydroxylation of indole rather than chlorohydroxylation should suppress isonitrile-derived byproduct formation, owing to the increased hardness of fluorine over chlorine. Therefore, a milder method to accomplish this transformation was developed, in which **21** was treated with a solution of XeF<sub>2</sub> in wet acetonitrile to provide **12** in 44% yield, *via* the intermediacy of **82**.<sup>14</sup> This reaction was routinely performed on more than 50 mg of **21**, and more than 580 mg of **12** has been prepared to date. A screen of various halogenating reagents confirmed that XeF<sub>2</sub> was the most efficient at promoting this reaction (Table 2). The chemoselectivity of this reagent is also noteworthy, given the presence of an olefin, which is known to react with XeF<sub>2</sub>,<sup>68</sup> and the reactive isonitrile moiety.

(67) The Wood group recently reported that their attempts to convert 10-*epi*-**78** into **12** were unsuccessful, utilizing a variety of conditions (see ref 17b).

Scheme 16. Synthesis of Isothiocyanate Derivatives<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a) DCM, CS(imid)<sub>2</sub> (3.3 equiv), 23 °C, 24 h, 60%; (b) THF, H<sub>2</sub>O, DDQ (2.5 equiv), 0 °C, 30 min, 68%; (c) DCM, XeF<sub>2</sub> (1.0 equiv), 23 °C, 15 min, 27%. DCM, dichloromethane; THF, tetrahydrofuran; DDQ, 2,3-dichloro-5,6-dicyanobenzoquinone.

Scheme 17. Synthesis and Degradation of 86<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a) DCM, TFDO (1.0 equiv), -78 °C, 5 min, 100%; (b) air, 23 °C, 6 d. DCM, dichloromethane; TFDO, methyl(trifluoromethyl)dioxirane.

In addition to **21** and **12** (*vide infra*), the isothiocyanate analogues can be readily synthesized *via* the sequence described herein (Scheme 16). Treatment of amine **70** with CS(imid)<sub>2</sub> provides isothiocyanate **83**, which can be oxidized to the 12-*epi*-fischerindole I analogue **84** (colorless cubes, mp 202 °C decomposition, see Scheme 16 for X-ray crystallographic analysis) in 76% yield. Treatment of **84** with XeF<sub>2</sub> provides the isothiocyanate analogue of welwitindolinone A (**85**) in an unoptimized 27% yield. Furthermore, the isocyanate analogue of welwitindolinone A (**86**) is directly available from welwit-

indolinone A (**12**) upon exposure to TFDO<sup>69</sup> (Scheme 17). However, this particular analogue exhibited greater instability than the isonitrile and isothiocyanate derivatives, and dimerized to the urea dimer **87** upon storage in air at room temperature. Despite this reactivity, it is conceivable that such intermediates might be useful for the preparation of further members of the welwitindolinone alkaloid family.

### Oxidation Attempts toward Welwitindolinone B

In order to test the biosynthetic hypothesis delineated in Scheme 2, and in an attempt to generate the welwitindolinone B core structure, welwitindolinone A (**12**) was subjected to a variety of epoxidation and general oxidation conditions. With the exception of TFDO that generated the corresponding isocyanate **86** (*vide supra*), all attempted conditions led to either recovered starting material or decomposition (see Supporting Information for a list of failed conditions). Consequently, three-step alternatives were envisioned, with the hope that the electronics of the tetrasubstituted olefin would be modified in a subtle fashion so as to permit oxidative rearrangement (see Scheme 18). Although not ideal, transformation of the isonitrile moiety into a *gem*-dibromide (such as **88**) or a formamide (such as **90**), oxidative rearrangement into the corresponding welwitindolinone B framework (**89** or **91**, respectively), followed by restoration of the original isonitrile group, could enable the formation of “welwitindolinone B isonitrile” (**14a**). With this strategy in mind, *gem*-dibromide **88** and formamide **90** were prepared from welwitindolinone A using phenyltrimethylammonium tribromide and formic acid, respectively, both in quantitative yields; functional group restoration to the original isonitrile **12** was also verified by using triethyl phosphite and phosgene, respectively. Unfortunately, all oxidation conditions experimented upon **88** or **90** resulted in either recovered starting material, welwitindolinone A (only when using **88**), or decomposition (see Supporting Information). Since some welwitindolinones are methylated at the indole nitrogen atom (see Chart 1), N(1)-methylformamide **92** was prepared from **12** in an attempt to alter reactivity, but this substrate also failed to undergo oxidative rearrangement (see Supporting Information).

These preliminary results do not necessarily contradict the biosynthetic hypothesis toward welwitindolinone B put forth in Scheme 2; perhaps the oxidation requisite for this transformation is enzymatically controlled.

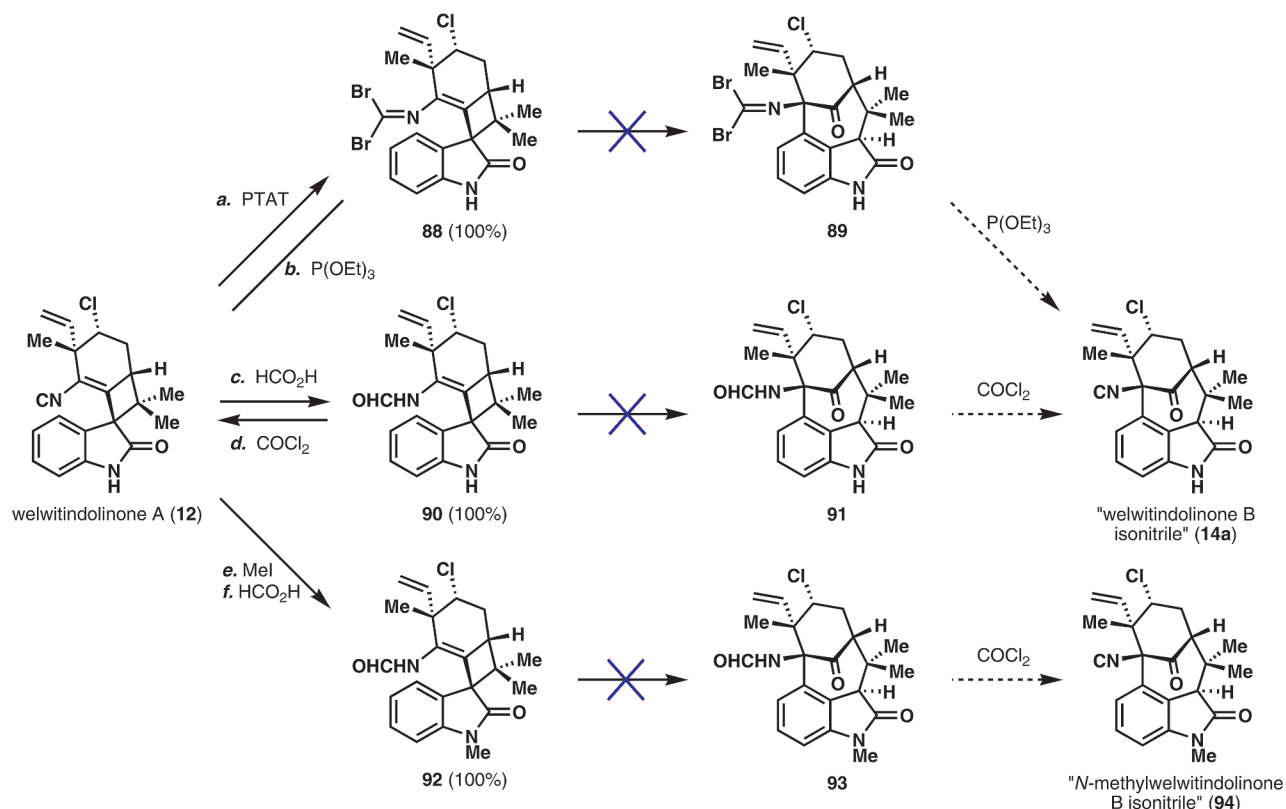
### Strategy Analysis and Conclusions

Due to their stunning molecular architectures and potent bioactivities, the hapalindoles, fischerindoles, and welwitindolinones were targeted for total synthesis, with the goals of inventing useful chemistry, discovering basic reactivity, understanding their biosynthesis, and allowing access to large quantities of these rare marine natural products. It is instructive to evaluate these syntheses from the vantage points of chemoselectivity, stereocontrol, and “redox economy”.

Numerous steps throughout this synthesis exhibited high levels of chemoselectivity, despite the presence of other reactive functionality. For example, the conversion of **46** to **45** proceeds in good yield in the presence of a chlorine atom that could potentially be eliminated under the reaction conditions. The vast majority of conditions screened for the cyclization of **45** to **44** formed numerous byproducts (including **65**, **66**, and **67**);

(68) Shellhamer, D. F.; Carter, D. L.; Chiacco, M. C.; Harris, T. E.; Henderson, R. D.; Low, W. S. C.; Metcalf, B. T.; Willis, M. C.; Hoesley, V. L.; Chapman, R. D. *J. Chem. Soc., Perkin Trans. 2* **1991**, 401–403.

(69) Mello, R.; Fiorentino, M.; Fusco, C.; Curci, R. *J. Am. Chem. Soc.* **1989**, *111*, 6749–6757.

Scheme 18. Oxidative Rearrangement Attempts<sup>a</sup>

<sup>a</sup> Reagents and conditions: (a) PTAT (1 equiv), DCM, -78 °C, 3 min, 100%; (b) P(OEt)<sub>3</sub>, DCM, 23 °C, 3 min, 100%; (c) HCO<sub>2</sub>H, H<sub>2</sub>O, THF, 4 °C, 12 h, 100%; (d) TEA, COCl<sub>2</sub>, DCM, 0 °C, 10 min, 82%; (e) MeI, K<sub>2</sub>CO<sub>3</sub>, acetone, 23 °C, 12 h, 100%; (f) HCO<sub>2</sub>H, H<sub>2</sub>O, THF, 4 °C, 12 h, 100%, PTAT, phenyltrimethylammonium tribromide; DCM, dichloromethane; THF, tetrahydrofuran; TEA, triethylamine.

Table 2. Ring Contraction Optimization

oxidant	yield (%)
<i>t</i> -BuOCl	0–25
NBS	0
Synfluor	0
Selectfluor	0
BMAST	0
NFSI	16
XeF <sub>2</sub>	44

however, the use of Montmorillonite K-10 acidic clay completely avoided such chemical entities. The conversion of **75** into **21** proceeds in excellent yield, despite the ease with which isonitriles can be oxidized, and the conversion of **21** to **12** proceeds in good yield, even though isonitriles have been observed to react with halogenating reagents.<sup>14</sup> Functional group manipulations were minimized, and the percentage of C–C bond-forming reactions was maximized. It is difficult to imagine how any steps could be “removed” from these syntheses, since all are necessary for the installation of requisite C–C bonds, functional groups, or key stereocenters. Furthermore, no protecting groups are utilized throughout the course of this entire synthesis, despite numerous opportunities in which they could

potentially have been employed to circumvent undesirable reactivity. In fact, rather than resorting to protecting group chemistry, the innate reactivities of the functional groups were employed, which led to the invention of new chemistry (direct indole coupling, extremely mild fluorohydroxylative ring-contraction using XeF<sub>2</sub>, installation of the key quaternary stereocenter and neopentyl chlorine atom) or discovery of intriguing reactivity (**49** to **50**, **45** to **65**, **66**, and **67**, **71** to **21** and **74**, **75** to **21**, **44** to **78**, and **21** to **12**).

High levels of stereochemical induction are observed throughout the synthesis. For example, the direct indole coupling reaction (**31** and **45**) provides a single diastereomer of coupled product, and the reductive alkylation of **31** to provide **41** gives only a single diastereomer. The reductive aminations proceed in moderate (**41** to **27**, which could be increased with the use of microwave irradiation) to complete diastereoselectivity (**43** to **29** and **44** to **70**). Furthermore, by utilizing XeF<sub>2</sub>, complete diastereomeric induction is observed in the conversion of **21** to **12**, even though the facial bias for this transformation is minimal.

Finally, the syntheses are characterized by an adherence to the concept of “redox economy”. Analogous to “atom economy”<sup>70</sup> or “step economy”,<sup>71</sup> “redox economy”<sup>72</sup> minimizes superfluous redox manipulations within a synthesis; rather,

(70) Trost, B. M. *Science* **1991**, *254*, 1471–1477.

(71) Wender, P. A.; Miller, B. L. In *Organic Synthesis: Theory and Applications*; Hudlicky, T., Ed.; JAI Press: Greenwich, CT, 1993, Vol. 2.

(72) For illuminating discussions of oxidation state control in synthesis, see: (a) Evans, D. A.; Andrews, G. C. *Acc. Chem. Res.* **1974**, *7*, 147–155. (b) Hendrickson, J. B. *J. Am. Chem. Soc.* **1975**, *97*, 5784–5800.



the oxidation state of intermediates linearly and steadily increases throughout the course of the synthesis. Only one reduction step (reductive amination) is utilized during these syntheses, which is strategically placed to install a key stereocenter. In fact, the flexible adherence of these syntheses to the proposed biogenesis of these alkaloids reinforces the minimization of redox reactions, as is commonly observed in Nature's biosynthesis of terpenes and alkaloids. Such considerations allowed the efficient, practical, and concise syntheses of numerous members of this natural product family (**10**, **12**, **21**, **27**, **29**, and **42**). Overall, this total synthesis program is characterized by inventive retrosynthetic disconnections, which rapidly assemble the skeletal structure and allow for rapid increase in complexity. These studies are yet another example of how natural products can catalyze new discoveries in chemical reactivity.

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**Note Added after ASAP Publication.** Due to a production error the graphics for Schemes 4, 10, 11, 13, 14, and 16 were incomplete. These errors have been corrected for the versions posted on December 3, 2008, and in print.

**Supporting Information Available:** Full characterization, including  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra, and experimental procedures for selected compounds; X-ray crystallographic data, in CIF format, for **31**, **35**, **45**, **67**, **74**, **78**, and **84**. This material is available free of charge *via* the Internet at <http://pubs.acs.org>.

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